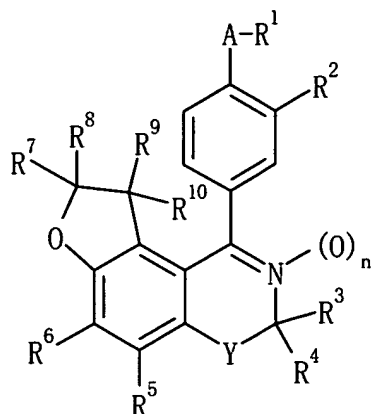


1. (ORIGINAL) A compound represented by the formula



wherein A represents (1) a bond, (2) a group represented by the formula $-\text{CR}^a=\text{CR}^b-$ (R^a and R^b each represent a hydrogen atom or a C_{1-6} alkyl group), (3) a group represented by the formula $-(\text{CONH})_p-(\text{C}(\text{R}^c)(\text{R}^d))_q-$ (R^c and R^d each represent a hydrogen atom or a C_{1-6} alkyl group, p represents 0 or 1 and q represents 1 or 2), (4) a group represented by the formula $-\text{CH}_2\text{OCH}_2-$ or (5) a group represented by the formula $-\text{OCH}_2-$;

R^1 represents (1) a cyano group or (2) an optionally esterified or amidated carboxyl group;

R^2 represents (1) a hydrogen atom, (2) an optionally substituted hydroxy group, (3) an optionally substituted amino group, (4) an optionally substituted alkyl group, (5) an optionally esterified or amidated carboxyl group or (6) a nitro group, or R^2 and A or R^1 may be taken together with the adjacent carbon atom to form a ring;

R^3 and R^4 each represent (1) a hydrogen atom, (2) an optionally substituted hydrocarbon group or (3) an acyl group, or

R^3 and R^4 may be taken together with the adjacent carbon atom to form an optionally substituted 3- to 8-membered ring;

R^5 represents (1) a hydrogen atom, (2) a cyano group, (3) an optionally substituted hydrocarbon group, (4) an acyl group or (5) an optionally substituted hydroxy group;

R^6 represents (1) a hydrogen atom, (2) an optionally substituted hydrocarbon group, (3) an acyl group, (4) an optionally substituted heterocyclic group, (5) a halogen atom, (6) an optionally substituted hydroxy group, (7) an optionally substituted thiol group, (8) a group represented by the formula $-S(O)_rR^{11}$ (R^{11} represents an optionally substituted hydrocarbon group or an optionally substituted heterocyclic group and r is 1 or 2) or (9) an optionally substituted amino group;

R^7 and R^8 each represent (1) a hydrogen atom or (2) an optionally substituted hydrocarbon group, or R^7 and R^8 may be taken together with the adjacent carbon atom to form an optionally substituted 3- to 8-membered ring;

R^9 and R^{10} each represent (1) a hydrogen atom or (2) an optionally substituted hydrocarbon group;

Y represents an optionally substituted methylene group; and n represents 0 or 1,

provided that if A is a bond, R^2 is not a hydrogen atom, and if A is a group represented by the formula $-(CONH)_p-(C(R^c)(R^d))_q-$ (R^c and R^d each represent a hydrogen atom or a C_{1-6} alkyl group, p represents 0 or 1 and q represents 1 or 2), R^6 is not methoxy, or a salt thereof.

2. (ORIGINAL) The compound according to claim 1, wherein R¹ is (i) a cyano group, (ii) a carboxyl group, (iii) a C₁₋₆ alkoxy-carbonyl group which may have 1 to 5 substituents selected from a group consisting of (1) a halogen atom, (2) a C₁₋₃ alkylenedioxy group, (3) a nitro group, (4) a cyano group, (5) an optionally halogenated C₁₋₆ alkyl group, (6) an optionally halogenated C₂₋₆ alkenyl group, (7) an optionally halogenated C₂₋₆ alkynyl group, (8) a C₃₋₈ cycloalkyl group, (9) a C₆₋₁₄ aryl group, (10) an optionally halogenated C₁₋₆ alkoxy group, (11) an optionally halogenated C₁₋₆ alkylthio group, (12) a hydroxy group, (13) an amino group, (14) a mono-C₁₋₆ alkylamino group, (15) a mono-C₆₋₁₄ arylamino group, (16) a di-C₁₋₆ alkylamino group, (17) a di-C₆₋₁₄ arylamino group, (18) an acyl group selected from formyl, carboxyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-

thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, sulfinio, sulfo, C₁₋₆ alkoxy-sulfinyl, C₆₋₁₄ aryloxy-sulfinyl, C₁₋₆ alkoxy-sulfonyl and C₆₋₁₄ aryloxy-sulfonyl, (19) an acylamino group selected from formylamino, C₁₋₆ alkyl-carboxamide, C₆₋₁₄ aryl-carboxamide, C₁₋₆ alkoxy-carboxamide, C₁₋₆ alkylsulfonylamino and C₆₋₁₄ arylsulfonylamino, (20) an acyloxy group selected from C₁₋₆ alkyl-carbonyloxy, C₆₋₁₄ aryl-carbonyloxy, C₁₋₆ alkoxy-carbonyloxy, mono-C₁₋₆ alkyl-carbamoyloxy, di-C₁₋₆ alkyl-carbamoyloxy, mono-C₆₋₁₄ aryl-carbamoyloxy, di-C₆₋₁₄ aryl-carbamoyloxy and nicotinoyloxy, (21) a 5- to 14-membered heterocyclic group containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, (22) a phosphono group, (23) a C₆₋₁₄ aryloxy group, (24) a di-C₁₋₆ alkoxy-phosphoryl group, (25) a C₆₋₁₄ arylthio group, (26) a hydrazino group, (27) an imino group, (28) an oxo group, (29) an ureido group, (30) a C₁₋₆ alkyl-ureido group, (31) a di-C₁₋₆ alkyl-ureido group, (32) an oxide

group and (33) a group formed by binding of 2 or 3 groups selected from (1) to (32) listed above and the like (hereinafter, abbreviated as Substituent group A), (iv) a C₃₋₈ cycloalkyloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (v) a C₇₋₁₆ aralkyloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (vi) a C₆₋₁₄ aryloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (vii) a carbamoyl group, (viii) a mono-C₁₋₆ alkyl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above, (ix) a di-C₁₋₆ alkyl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above, (x) a mono-C₆₋₁₄ aryl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above or (xi) a di-C₆₋₁₄ aryl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above,

R² is (i) a hydrogen atom, (ii) a group represented by the formula -OR¹² (R¹² represents (a) a hydrogen atom, (b) a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, or (c) an acyl group selected from formyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-

carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, C₁₋₆ alkoxysulfinyl, C₆₋₁₄ aryloxysulfinyl, C₁₋₆ alkoxysulfonyl and C₆₋₁₄ aryloxysulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above), (iii) a group represented by the formula -NR¹³R¹⁴ (R¹³ and R¹⁴ are each (i') a hydrogen atom, (ii') a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or

C_{7-16} aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, (iii') an acyl group selected from formyl, carbamoyl, C_{1-6} alkyl-carbonyl, C_{3-8} cycloalkyl-carbonyl, C_{1-6} alkoxy-carbonyl, C_{6-14} aryl-carbonyl, C_{7-16} aralkyl-carbonyl, C_{6-14} aryloxy-carbonyl, C_{7-16} aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono- C_{1-6} alkyl-carbamoyl, di- C_{1-6} alkyl-carbamoyl, mono- C_{6-14} aryl-carbamoyl, di- C_{6-14} aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C_{1-6} alkyl-thiocarbonyl, C_{3-8} cycloalkyl-thiocarbonyl, C_{1-6} alkoxy-thiocarbonyl, C_{6-14} aryl-thiocarbonyl, C_{7-16} aralkyl-thiocarbonyl, C_{6-14} aryloxy-thiocarbonyl, C_{7-16} aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono- C_{1-6} alkyl-thiocarbamoyl, di- C_{1-6} alkyl-thiocarbamoyl, mono- C_{6-14} aryl-thiocarbamoyl, di- C_{6-14} aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono- C_{1-6} alkylsulfamoyl, di- C_{1-6} alkylsulfamoyl, C_{6-14} arylsulfamoyl, C_{1-6} alkylsulfonyl, C_{6-14} arylsulfonyl, C_{1-6} alkylsulfinyl, C_{6-14} arylsulfinyl, C_{1-6} alkoxysulfinyl, C_{6-14} aryloxysulfinyl, C_{1-6} alkoxysulfonyl and C_{6-14}

aryloxysulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above or (iv') a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above, or R^{13} and R^{14} may be taken together with the adjacent a nitrogen atom to form a 5- to 14-membered ring), (iv) a C_{1-6} alkylideneamino group which may have 1 to 5 substituents selected from Substituent group A described above, (v) a C_{1-6} alkyl group which may have 1 to 5 substituents selected from Substituent group A described above, (vi) a carboxyl group, (vii) a C_{1-6} alkoxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (viii) a C_{3-8} cycloalkyloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (ix) a C_{7-16} aralkyloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (x) a C_{6-14} aryloxy-carbonyl group which may have 1 to 5 substituents selected from Substituent group A described above, (xi) a carbamoyl group, (xii) a mono- C_{1-6} alkyl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above, (xiii) a di- C_{1-6} alkyl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above, (xiv) a mono- C_{6-14} aryl-carbamoyl group which may have 1 to 5 substituents selected from Substituent group A described above, (xv) a di- C_{6-14} aryl-carbamoyl group

which may have 1 to 5 substituents selected from Substituent group A described above or (xvi) a nitro group, or R^2 and A or R^1 may be taken together to form a 5- to 14-membered ring containing 1 to 4 hetero atoms selected from a nitrogen atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above;

each of R^3 and R^4 is any of the following (i) to (iii):

(i) a hydrogen atom,

(ii) a C_{1-6} alkyl group, C_{2-6} alkenyl group, C_{2-6} alkynyl group, C_{3-8} cycloalkyl group, C_{3-8} cycloalkenyl group, C_{6-14} aryl group or C_{7-16} aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above,

(iii) an acyl group selected from formyl, carboxyl, carbamoyl, C_{1-6} alkyl-carbonyl, C_{3-8} cycloalkyl-carbonyl, C_{1-6} alkoxy-carbonyl, C_{6-14} aryl-carbonyl, C_{7-16} aralkyl-carbonyl, C_{6-14} aryloxy-carbonyl, C_{7-16} aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono- C_{1-6} alkyl-carbamoyl, di- C_{1-6} alkyl-carbamoyl, mono- C_{6-14} aryl-carbamoyl, di- C_{6-14} aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C_{1-6} alkyl-thiocarbonyl, C_{3-8} cycloalkyl-thiocarbonyl, C_{1-6} alkoxy-thiocarbonyl, C_{6-14} aryl-thiocarbonyl, C_{7-16} aralkyl-thiocarbonyl, C_{6-14} aryloxy-thiocarbonyl, C_{7-16} aralkyloxy-thiocarbonyl, 5- or 6-membered

heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, sulfinio, sulfo, C₁₋₆ alkoxy sulfinyl, C₆₋₁₄ aryloxy sulfinyl, C₁₋₆ alkoxy sulfonyl and C₆₋₁₄ aryloxy sulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above; or

R³ and R⁴ may be taken together with the adjacent carbon atom to form C₃₋₈ cycloalkane or a 3- to 8-membered heterocycle, which may have respectively 1 to 3 substituents selected from C₁₋₆ alkyl, C₆₋₁₄ aryl, C₇₋₁₆ aralkyl, amino, mono-C₁₋₆ alkylamino, mono-C₆₋₁₄ arylamino, di-C₁₋₆ alkylamino, di-C₆₋₁₄ arylamino and a 4- to 10-membered aromatic heterocyclic group,

R⁵ is (i) a hydrogen atom, (ii) a cyano group, (iii) a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, (iv) an acyl group selected from formyl, carboxyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈

cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, sulfinio, sulfo, C₁₋₆ alkoxy-sulfinyl, C₆₋₁₄ aryloxy-sulfinyl, C₁₋₆ alkoxy-sulfonyl and C₆₋₁₄ aryloxy-sulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above, or (v) a group represented by the formula -OR¹⁵ (R¹⁵ represents

(a) a hydrogen atom, (b) a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, or (c) an acyl group selected from formyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄

arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, C₁₋₆ alkoxyulfinyl, C₆₋₁₄ aryloxyulfinyl, C₁₋₆ alkoxyulfonyl and C₆₋₁₄ aryloxyulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above),

R⁶ is any of the following (i) to (x):

(i) a hydrogen atom,

(ii) a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above,

(iii) an acyl group selected from formyl, carboxyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected

from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, sulfinio, sulfo, C₁₋₆ alkoxy sulfinyl, C₆₋₁₄ aryloxy sulfinyl, C₁₋₆ alkoxy sulfonyl and C₆₋₁₄ aryloxy sulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above,

(iv) a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above,

(v) a halogen atom,

(vi) a group represented by the formula -OR¹⁶ (R¹⁶ represents (i') a hydrogen atom, (ii') a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, (iii') an acyl group selected from formyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl

containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, C₁₋₆ alkoxysulfinyl, C₆₋₁₄ aryloxysulfinyl, C₁₋₆ alkoxysulfonyl and C₆₋₁₄ aryloxysulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above, or (iv') a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which

may have 1 to 5 substituents selected from Substituent group A described above),

(vii) a group represented by the formula $-SR^{17}$ (R^{17} represents (i') a hydrogen atom, (ii') a C_{1-6} alkyl group, C_{2-6} alkenyl group, C_{2-6} alkynyl group, C_{3-8} cycloalkyl group, C_{3-8} cycloalkenyl group, C_{6-14} aryl group or C_{7-16} aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, (iii') an acyl group selected from formyl, carbamoyl, C_{1-6} alkyl-carbonyl, C_{3-8} cycloalkyl-carbonyl, C_{1-6} alkoxy-carbonyl, C_{6-14} aryl-carbonyl, C_{7-16} aralkyl-carbonyl, C_{6-14} aryloxy-carbonyl, C_{7-16} aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono- C_{1-6} alkyl-carbamoyl, di- C_{1-6} alkyl-carbamoyl, mono- C_{6-14} aryl-carbamoyl, di- C_{6-14} aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C_{1-6} alkyl-thiocarbonyl, C_{3-8} cycloalkyl-thiocarbonyl, C_{1-6} alkoxy-thiocarbonyl, C_{6-14} aryl-thiocarbonyl, C_{7-16} aralkyl-thiocarbonyl, C_{6-14} aryloxy-thiocarbonyl, C_{7-16} aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono- C_{1-6} alkyl-thiocarbamoyl, di- C_{1-6} alkyl-thiocarbamoyl, mono- C_{6-14} aryl-thiocarbamoyl, di- C_{6-14} aryl-thiocarbamoyl, 5- or 6-membered

heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, C₁₋₆ alkoxy sulfinyl, C₆₋₁₄ aryloxy sulfinyl, C₁₋₆ alkoxy sulfonyl and C₆₋₁₄ aryloxy sulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above or (iv') a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above),

(viii) a group represented by the formula $-S(O)_r R^{11}$ (R^{11} represents (i') a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above or (ii') a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above and r is 1 or 2) or

(ix) a group represented by the formula $-NR^{18}R^{19}$ (R^{18} and R^{19} each represent (i') a hydrogen atom, (ii') a C₁₋₆ alkyl group, C₂₋₆ alkenyl group, C₂₋₆ alkynyl group, C₃₋₈ cycloalkyl group, C₃₋₈ cycloalkenyl group, C₆₋₁₄ aryl group or C₇₋₁₆ aralkyl group, each

of which may have 1 to 5 substituents selected from Substituent group A described above, (iii') an acyl group selected from formyl, carbamoyl, C₁₋₆ alkyl-carbonyl, C₃₋₈ cycloalkyl-carbonyl, C₁₋₆ alkoxy-carbonyl, C₆₋₁₄ aryl-carbonyl, C₇₋₁₆ aralkyl-carbonyl, C₆₋₁₄ aryloxy-carbonyl, C₇₋₁₆ aralkyloxy-carbonyl, 5- or 6-membered heterocyclic carbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, mono-C₁₋₆ alkyl-carbamoyl, di-C₁₋₆ alkyl-carbamoyl, mono-C₆₋₁₄ aryl-carbamoyl, di-C₆₋₁₄ aryl-carbamoyl, 5- or 6-membered heterocyclic carbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, C₁₋₆ alkyl-thiocarbonyl, C₃₋₈ cycloalkyl-thiocarbonyl, C₁₋₆ alkoxy-thiocarbonyl, C₆₋₁₄ aryl-thiocarbonyl, C₇₋₁₆ aralkyl-thiocarbonyl, C₆₋₁₄ aryloxy-thiocarbonyl, C₇₋₁₆ aralkyloxy-thiocarbonyl, 5- or 6-membered heterocyclic thiocarbonyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, thiocarbamoyl, mono-C₁₋₆ alkyl-thiocarbamoyl, di-C₁₋₆ alkyl-thiocarbamoyl, mono-C₆₋₁₄ aryl-thiocarbamoyl, di-C₆₋₁₄ aryl-thiocarbamoyl, 5- or 6-membered heterocyclic thiocarbamoyl containing 1 to 3 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, sulfamoyl, mono-C₁₋₆ alkylsulfamoyl, di-C₁₋₆ alkylsulfamoyl, C₆₋₁₄ arylsulfamoyl, C₁₋₆ alkylsulfonyl, C₆₋₁₄ arylsulfonyl, C₁₋₆ alkylsulfinyl, C₆₋₁₄ arylsulfinyl, C₁₋₆ alkoxysulfinyl, C₆₋₁₄ aryloxysulfinyl, C₁₋₆ alkoxysulfonyl and C₆₋₁₄

aryloxysulfonyl, which may have 1 to 5 substituents selected from Substituent group A described above or (iv') a 5- to 14-membered heterocycle containing 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may have 1 to 5 substituents selected from Substituent group A described above),

R^7 and R^8 are each (i) a hydrogen atom or (ii) a C_{1-6} alkyl group, C_{2-6} alkenyl group, C_{2-6} alkynyl group, C_{3-8} cycloalkyl group, C_{3-8} cycloalkenyl group, C_{6-14} aryl group or C_{7-16} aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, or R^7 and R^8 may be taken together with the adjacent carbon atom to form C_{3-8} cycloalkane or a 3- to 8-membered heterocycle, which may have respectively 1 to 3 substituents selected from C_{1-6} alkyl, C_{6-14} aryl, C_{7-16} aralkyl, amino, mono- C_{1-6} alkylamino, mono- C_{6-14} arylamino, di- C_{1-6} alkylamino, di- C_{6-14} arylamino and a 4- to 10-membered aromatic heterocyclic group;

R^9 and R^{10} are each (i) a hydrogen atom or (ii) a C_{1-6} alkyl group, C_{2-6} alkenyl group, C_{2-6} alkynyl group, C_{3-8} cycloalkyl group, C_{3-8} cycloalkenyl group, C_{6-14} aryl group or C_{7-16} aralkyl group, each of which may have 1 to 5 substituents selected from Substituent group A described above, and

Y is a methylene group which may have 1 or 2 substituents selected from Substituent group A described above.

3. (ORIGINAL) The compound according to claim 1, wherein A is (1) a bond, (2) a group represented by the formula $-CR^a=CR^b-$ (R^a and R^b each represent a hydrogen atom or a C_{1-6} alkyl group), (3) a group represented by the formula $-(CONH)_p-(C(R^c)(R^d))_q-$ (R^c and R^d each represent a hydrogen atom or a C_{1-6} alkyl group, p represents 0 or 1 and q represent s 1 or 2), (4) a group represented by the formula $-CH_2OCH_2-$ or (5) a group represented by the formula $-OCH_2-$,

R^1 is (1) a cyano group, (2) a carboxyl group, (3) a C_{1-6} alkoxy carbonyl group, (4) a carbamoyl group or (5) an N-mono- C_{1-6} alkyl carbamoyl group,

R^2 is (1) a hydrogen atom, (2) a hydroxy group, (3) a C_{1-6} alkoxy group, (4) a C_{7-16} aralkyloxy group, (5) an amino group, (6) a mono- C_{1-6} alkylamino group which may have one substituent selected from carboxyl, carbamoyl, quinolyl, phenoxy and pyridyl, (7) a mono- C_{7-16} aralkylamino group which may have one substituent selected from a halogen atom, cyano, C_{1-6} alkoxy, carboxyl and C_{1-6} alkoxy carbonyl, (8) a mono- C_{6-14} arylamino group, (9) a mono- C_{1-6} alkyl carbonylamino group which may have 1 to 3 substituents selected from a halogen atom, thienyl and C_{1-6} alkoxy carbonyl- C_{1-6} alkylthio, (10) a mono- C_{1-6} alkylsulfonylamino group, (11) a mono- C_{6-14} aryl carbonylamino group which may have one substituent selected from C_{1-6} alkoxy and C_{1-6} alkyl carbonylamino, (12) a quinolyl carbonylamino group, (13) a pyridyl carbonylamino group which may have 1 or 2 halogen atoms, (14) an indolyl carbonylamino group, (15) a N- C_{1-6} alkyl-N- C_{1-6} alkyl carbonylamino group which

may have 1 to 4 substituents selected from a halogen atom, C₁₋₆ alkoxy-carbonyl and quinolyl, (16) a N-C₁₋₆ alkylcarbonyl-N-C₇₋₁₆ aralkylamino group which may have 1 to 3 halogens, (17) a N-C₁₋₆ alkyl-N-pyridylcarbonylamino group, (18) a C₁₋₆ alkylideneamino group which may have one di-C₁₋₆ alkylamino, (19) a mono-C₁₋₆ alkylureido group which may have one C₁₋₆ alkoxy-carbonyl, (20) a di-C₁₋₆ alkylureido, (21) a mono-C₆₋₁₄ arylureido group, (22) a 1-imidazolidinyl group which may have 1 to 3 substituents selected from C₁₋₆ alkyl and oxo, (23) a C₁₋₆ alkyl group, (24) a C₁₋₆ alkoxy-carbonyl group, (25) a nitro group or (26) a 1-pyrrolidinyl group, or

R² and A or R¹ may be taken together with the adjacent carbon atom to form a nitrogen-containing 5- to 7-membered ring which may have 1 to 3 substituents selected from (1) a hydroxy group, (2) C₁₋₆ alkyl which may have one C₁₋₆ alkoxy-carbonyl, (3) C₇₋₁₆ aralkyl, (4) C₆₋₁₄ aryl and (5) oxo,

R³ and R⁴ are each a C₁₋₆ alkyl group,

R⁵ is a hydrogen atom,

R⁶ is a C₁₋₆ alkoxy group,

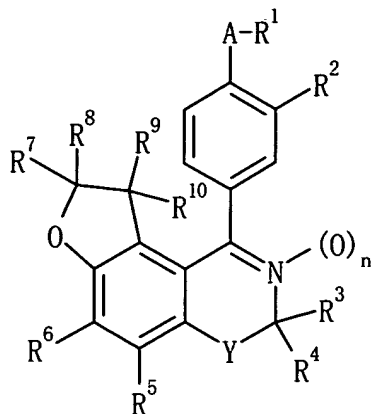
R⁷ and R⁸ are each a C₁₋₆ alkyl group,

R⁹ and R¹⁰ are each a hydrogen atom,

Y is a methylene group and

n is 0.

4. (ORIGINAL) A compound represented by the formula



wherein A is (1) a bond, (2) a group represented by the formula $-\text{CH}=\text{CH}-$, (3) a group represented by the formula $-\text{CONH}-\text{C}(\text{R}^c)(\text{R}^d)-$ (R^c and R^d are each a hydrogen atom or a C_{1-6} alkyl group), or (4) a group represented by the formula $-\text{OCH}_2-$,

R^1 is (1) a cyano group, (2) a carboxyl group, (3) a C_{1-6} alkoxy carbonyl group, (4) a carbamoyl group or (5) an N-mono- C_{1-6} alkyl carbamoyl group,

R^2 is (1) a hydroxy group, (2) a C_{1-6} alkoxy group, (3) a C_{7-16} aralkyloxy group, (4) an amino group, (5) a mono- C_{1-6} alkylamino group which may have one substituent selected from carboxyl, carbamoyl, quinolyl, phenoxy and pyridyl, (6) a mono- C_{7-16} aralkylamino group which may have one substituent selected from a halogen atom, cyano, C_{1-6} alkoxy, carboxyl and C_{1-6} alkoxy carbonyl, (7) a mono- C_{6-14} arylamino group, (8) a mono- C_{1-6} alkyl carbonylamino group which may have 1 to 3 substituents selected from a halogen atom, thienyl and C_{1-6} alkoxy carbonyl- C_{1-6} alkylthio, (9) a mono- C_{1-6} alkylsulfonylamino group, (10) a mono- C_{6-14} aryl carbonylamino group which may have one substituent selected from C_{1-6} alkoxy and C_{1-6} alkyl carbonylamino, (11) a

quinolylcarbonylamino group, (12) a pyridylcarbonylamino group which may have 1 or 2 halogen atoms, (13) an indolylcarbonylamino group, (14) a N-C₁₋₆ alkyl-N-C₁₋₆ alkylcarbonylamino group which may have 1 to 4 substituents selected from a halogen atom, C₁₋₆ alkoxy-carbonyl and quinolyl, (15) a N-C₁₋₆ alkylcarbonyl-N-C₇₋₁₆ aralkylamino group which may have 1 to 3 halogens, (16) a N-C₁₋₆ alkyl-N-pyridylcarbonylamino group, (17) a C₁₋₆ alkylideneamino group which may have one di-C₁₋₆ alkylamino, (18) a mono-C₁₋₆ alkylureido group which may have one C₁₋₆ alkoxy-carbonyl, (19) a di-C₁₋₆ alkylureido group, (20) a mono-C₆₋₁₄ arylureido group, (21) a 1-imidazolidinyl group which may have 1 to 3 substituents selected from C₁₋₆ alkyl and oxo, (22) a C₁₋₆ alkyl group, (23) a C₁₋₆ alkoxy-carbonyl group, (24) a nitro group or (25) a 1-pyrrolidinyl group, or R² and A or R¹ may be taken together with the adjacent carbon atom to form a nitrogen-containing 5- to 7-membered ring which may have 1 to 3 substituents selected from (1) a hydroxy group, (2) a C₁₋₆ alkyl group which may have one C₁₋₆ alkoxy-carbonyl, (3) a C₇₋₁₆ aralkyl group, (4) a C₆₋₁₄ aryl group and (5) an oxo group,

R³ and R⁴ are each a C₁₋₆ alkyl group,

R⁵ is a hydrogen atom,

R⁶ is a C₂₋₆ alkoxy group,

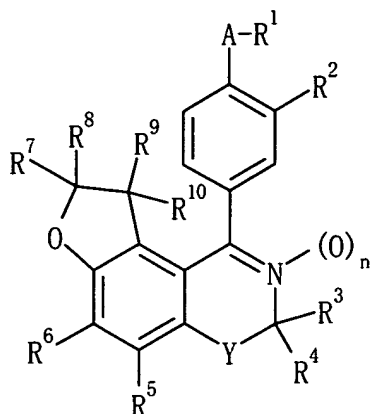
R⁷ and R⁸ are each a C₁₋₆ alkyl group,

R⁹ and R¹⁰ are each a hydrogen atom,

Y is a methylene group, and

n is 0, or a salt thereof.

5. (ORIGINAL) A compound represented by the formula



wherein A is (1) a group represented by the formula -
 $\text{CR}^{\text{a}}=\text{CR}^{\text{b}}-$ (R^{a} and R^{b} are each a hydrogen atom or a C_{1-6} alkyl
 group), (2) a group represented by the formula $-(\text{CONH})_{\text{p}}-$
 $(\text{C}(\text{R}^{\text{c}})(\text{R}^{\text{d}}))_{\text{q}}-$ (R^{c} and R^{d} are each a hydrogen atom or a C_{1-6} alkyl
 group, p is 0 or 1 and q is 1 or 2), (3) a group represented by
 the formula $-\text{CH}_2\text{OCH}_2-$ or (4) a group represented by the formula -
 OCH_2- ,

R^1 is (1) a carboxyl group, (2) a C_{1-6} alkoxy carbonyl group,
 (3) an N-mono- C_{1-6} alkyl carbamoyl group or (4) a carbamoyl group,

R^2 is a hydrogen atom,

R^3 and R^4 are each a C_{1-6} alkyl group,

R^5 is a hydrogen atom,

R^6 is a C_{2-6} alkoxy group,

R^7 and R^8 are each a C_{1-6} alkyl group,

R^9 and R^{10} are each a hydrogen atom,

Y is a methylene group, and

n is 0, or a salt thereof.

6. (ORIGINAL) The compound according to claim 4, wherein A is (1) a bond or (2) a group represented by the formula $-\text{CH}=\text{CH}-$.

7. (ORIGINAL) The compound according to claim 5, wherein A is (1) a group represented by the formula $-\text{CH}=\text{CH}-$, (2) a group represented by the formula $-(\text{C}(\text{R}^{\text{c}})(\text{R}^{\text{d}}))-$ (R^{c} and R^{d} each represent a hydrogen atom or a C_{1-6} alkyl group) or (3) a group represented by the formula $-\text{CH}_2\text{OCH}_2-$.

8. (ORIGINAL) The compound according to claim 4, wherein R^1 is a carboxyl group or a carbamoyl group.

9. (ORIGINAL) The compound according to claim 5, wherein R^1 is a carboxyl group.

10. (ORIGINAL) The compound according to claim 4, wherein R^2 is (1) a C_{1-6} alkoxy group, (2) a mono- C_{1-6} alkylamino group, (3) a mono- C_{7-16} aralkylamino group, (4) a quinolylcarbonylamino group or (5) a pyridylcarbonylamino group.

11. (CURRENTLY AMENDED) The compound according to claim 4 ~~or 5~~, wherein R^3 and R^4 are each methyl.

12. (CURRENTLY AMENDED) The compound according to claim 4 ~~or 5~~, wherein R^6 is ethoxy.

13. (CURRENTLY AMENDED) The compound according to claim 4 ~~or 5~~, wherein R⁷ and R⁸ are each methyl.

14. (ORIGINAL) The compound according to claim 4, which is 4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)-2-[(phenylmethyl)amino]benzoic acid, 4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)-2-(ethylamino)benzoic acid, (E)-3-[4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)-2-methoxyphenyl]-2-propenoic acid, 4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)-2-[(2-quinolinylcarbonyl)amino]benzoic acid, 4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)-2-[(2-pyridinylcarbonyl)amino]benzene acetic acid, N-[2-(aminocarbonyl)-5-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)phenyl]-2-pyridinecarboxamide or a salt thereof.

15. (ORIGINAL) The compound according to claim 5, which is [[4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)phenyl]methoxy]acetic acid, 4-(6-ethoxy-3,4,8,9-tetrahydro-3,3,8,8-tetramethylfuro[2,3-h]isoquinolin-1-yl)- α,α -dimethylbenzene acetic acid or a salt thereof.

16. (CURRENTLY AMENDED) ~~The~~ A pharmaceutical composition comprising the compound according to claim 1 or a prodrug thereof and a pharmaceutically acceptable carrier, excipient or diluent.

17. (ORIGINAL) The pharmaceutical composition according to claim 16, which is a phosphodiesterase IV inhibitor.

18. (ORIGINAL) The pharmaceutical composition according to claim 16, which is a prophylactic and/or therapeutic agent against inflammatory diseases, atopic dermatitis, allergic rhinitis, asthma, chronic obstructive pulmonary diseases, chronic rheumatoid arthritis, autoimmune diseases, depression, Alzheimer's dementia, memory disorders, osteoporosis, diabetes or atherosclerosis.

19. (ORIGINAL) A method of inhibiting phosphodiesterase IV, which comprises administering to a mammal an effective amount of the compound according to claim 1 or a prodrug thereof.

20. (ORIGINAL) A method of preventing and/or treating inflammatory diseases, atopic dermatitis, allergic rhinitis, asthma, chronic obstructive pulmonary diseases, chronic rheumatoid arthritis, autoimmune diseases, depression, Alzheimer's dementia, memory disorders, osteoporosis, diabetes or atherosclerosis, which comprises administering to a mammal an

effective amount of the compound according to claim 1 or a prodrug thereof.

21. (CURRENTLY AMENDED) A method for making a phosphodiesterase IV inhibitor pharmaceutical composition, said method comprising combining ~~Use of~~ the compound according to claim 1 or a prodrug thereof ~~for manufacturing a phosphodiesterase IV inhibitor~~ with a pharmaceutically acceptable carrier, excipient or diluent.

22. (CURRENTLY AMENDED) A method for making ~~Use of the compound according to claim 1 or a prodrug thereof for manufacturing~~ a prophylactic and/or therapeutic agent against inflammatory diseases, atopic dermatitis, allergic rhinitis, asthma, chronic obstructive pulmonary diseases, chronic rheumatoid arthritis, autoimmune diseases, depression, Alzheimer's dementia, memory disorders, osteoporosis, diabetes or atherosclerosis, said method comprising combining the compound according to claim 1 or a prodrug thereof with a pharmaceutically acceptable carrier, excipient or diluent.

23. (NEW) The compound according to claim 5, wherein R^3 and R^4 are each methyl.

24. (NEW) The compound according to claim 5, wherein R^6 is ethoxy.

25. (NEW) The compound according to claim 5, wherein R^7 and R^8 are each methyl.